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OM nucleic - nucleic search, using sw model

July 15, 2006, 18:04:58; Search time 547 Seconds Run on: (without alignments)

7647.805 Million cell updates/sec

Title: US-10-722-939-1-T39977 39901-40500

Perfect score:

Sequence: 1 tacaatagaccctgcttctt.....aataagtgacagagctgtga 600

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 5244920 segs, 3486124231 residues

Total number of hits satisfying chosen parameters: 10489840

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

Database : N_Geneseq_8:*

1: geneseqn1980s:*

2: geneseqn1990s:*

3: geneseqn2000s:*

4: geneseqn2001as:*

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14: geneseqn2005s:*

15: geneseqn2006s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB	ID	Description
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C	23 ′				4 14	AAS46704	
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C	30			110000 16766	5 6	AAI61373_0	Aai61373 Soybean 3
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PA
     (SEQU-) SEQUENOM INC.
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     Roth RB, Nelson MR, Braun A, Kammerer SM, Reneland R;
XX
DR
     WPI; 2004-441037/41.
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PT
     Identifying a subject at risk of breast cancer by detecting the presence
PT
     of polymorphic variations in the DLG1, KIAA0783, DPF3 or CENPC1 regions
PT
     which are associated with breast cancer in a nucleic acid sample from a
PT
     subject.
XX
PS
     Claim 24; Fig 1; 227pp; English.
XX
CC
     The present invention relates to a method for identifying a subject at
```

CC risk of breast cancer. The method comprising detecting the presence or CC absence of one or more polymorphic variations associated with breast CC cancer in a nucleic acid sample from a subject. The nucleic acid sample CC comprises the DLG1 region (ADO79402), KIAA0783 region (ADO79403), DPF3 CC region (AD079404) or CENPC1 region (AD079405). The gene DLGI (discs, CC large homolog 1 (Drosophila)) is also known as synapse-associated protein CC 97, hdlg or SAP97. DLG1 has been mapped to chromosomal position 3g29. The CC gene KIAA0783 is also known as PHF14 and PHD finger protein 14. KIAA0783 CC has been mapped to chromosomal position 7p21.3. The KIAA0783 protein is a CC novel gene with unknown function, however, being a zinc finger protein, CC it likely to be a transcription factor. The gene DPF3 (D4, zinc and CC double PHD fingers, family 3) is also known as CERD4, cer-d4, FLJ14079 CC and 2810403B03Rik. DPF3 is a Rho family guanine-nucleotide exchange CC factor. DPF3 has been mapped to chromosomal position 14q24.3-q31.1. The CC gene CENPCI (centromere protein C1) is also known as Centromere CC autoantigen C1. CENPC1 has been mapped to chromosomal position 4q12-CC q13.3. CENPC1 is a centromere autoantigen and a component of the inner CC kinetochore plate. The CENPC1 protein is required for maintaining proper CC kinetochore size and a timely transition to anaphase. The method is CC useful for identifying a subject at risk of breast cancer, for early diagnosis, prevention and treatment of breast cancer, to analyze and CC CC predict a response to a breast cancer treatment, and in clinical drug CC trials. XX SO Sequence 76600 BP; 25920 A; 14119 C; 13638 G; 22902 T; 0 U; 21 Other; 100.0%; Score 600; DB 12; Length 76600;

99.8%; Pred. No. 2.4e-84;

Query Match

Db

Best Local Similarity

Matches 599; Conservative 1; Mismatches 0; Indels 1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60 QУ Db 39901 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 39960 Qy 61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCCTGAGAAAACAATTAAATATTAA 120 39961 ATCTACTTCATTTACTYGTAATATACAGTCATTGACCCTGAGAAAACAATTAAATATTAA 40020 Db Qу Db QУ 181 CTTTTTTAAGGTAAAGAGAATTATAAATAATTCTGGAGTAATTCCAGAAAACATAAATGA 240 Db 40081 CTTTTTTAAGGTAAAGAGAATTATAAATAATTCTGGAGTAATTCCAGAAAACATAAATGA 40140 241 AGAAAGTATATCAAAAACTAATATAAACAAATACAAACATTTCCCAAGGGCCAGCAAAAG 300 Qу Db 40141 AGAAAGTATATCAAAAACTAATATAAACAAATACAAACATTTCCCAAGGGCCAGCAAAAG 40200 Qу Db Qу 361 ACCATTTGCATACTTCCTGCATGCTTATTCTGTGTAGGCATTAAGACACACTTTATAAAA 420

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XX
PA
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PΙ
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DR
    P-PSDB; AEE84581.
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PT
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PT
    inhibiting apoptosis for treating cancer or neurodegenerative diseases.
PT
    comprises semaphorin 4B protein and discs large homolog 1 and/or 3
PT
    protein.
XX
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Disclosure; SEQ ID NO 28; 140pp; Japanese.

PS XX CC

The invention relates to a complex (I) comprising semaphorin 4B (SEMA4B) protein (P1) having an amino acid sequence which is similar or substantially similar to that of (SEQ ID NO:26; see AEE84580), and discs large homolog 1 (DLG1) and/or DLG3 (P2) having an amino acid sequence which is similar or substantially similar to that of (SEQ ID N0:27 or SEQ ID NO:29; see AEE84581and AEE84583). (P1) of (I) further comprises one of four fully amino acid sequences (SEQ ID No. 1, 4, 7 or 10; see AEE84555, AEE84558, AEE84561 and AEE84564). Also included are: an antibody (II) with respect to (I), or that inhibits or promotes dissociation or formation of (I); a pharmaceutical (III) comprising (II); a diagnostic (IV) comprising (II); screening (M1) compound or its salt that inhibits or promotes binding of (P1) with (P2) or dissociation of (I), involves utilizing (P1) or (P2); a kit for carrying out (M1), comprising (P1) or (P2); a compound (V) or its salt that inhibits binding of (P1) with (P2) or promotes dissociation of (I); a promoter (VI) of apoptosis of cancer cell, the cancer proliferation inhibitor (VII) or preventive/therapeutic agent of cancer (VIII), comprising (V) or its salt; a compound (IX) or its salt that promotes binding of (P1) with (P2) or inhibits dissociation of (I); an inhibitor (X) of apoptosis of neuron, or preventive/therapeutic agent (XI) of neurodegenerative diseases, comprising (IX) or its salt; promoting (M2) apoptosis or inhibiting (M3) cancer cell proliferation, involves inhibiting binding of (P1) and (P2) or promoting dissociation of (I); preventing or treating (M4) cancer, involves inhibiting binding of (P1) and (P2) or promoting dissociation of (I); inhibiting (M5) apoptosis of neuron or preventing or treating (M6) neurodegenerative diseases, involves promoting binding of (P1) and (P2) or inhibiting dissociation of (I); screening compound or its salt for preventing or treating cancer or neurodegenerative diseases, involves utilizing (P2), its partial peptide or its salt; the use of a substance (XII) or its salt that inhibits binding of (P1) and (P2) or promotes dissociation of (I) for producing promoter of apoptosis of cancer cell, cancer cell proliferation inhibitor or preventive/therapeutic agent of cancer; and the use of a substance (XIII) or its salt that promotes binding of (P1) and (P2) or inhibits dissociation of (I) for producing inhibitor of apoptosis of neuron or preventive/therapeutic agent of neurodegenerative diseases. (III) is useful as promoter of apoptosis of cancer cell, cancer cell proliferation inhibitor or preventive/therapeutic agent of cancer, or as apoptosis inhibitor of neuron or preventive/therapeutic agent of neurodegenerative diseases. (IV) is useful for diagnosing cancer or neurodegenerative diseases. (VI), (VII) or (VIII) is useful for promoting apoptosis of cancer cell, inhibiting cancer proliferation, or preventing or treating cancer. (X) or (XI) is useful for inhibiting apoptosis of neuron, or preventing or treating neurodegenerative diseases. (M2) is useful for promoting apoptosis. (M3) is useful for inhibiting cancer cell proliferation. (M4) is useful for preventing or treating cancer. (M5) is useful for inhibiting apoptosis of neurons. (M6) is useful for preventing or treating neurodegenerative diseases. (XII) is useful for producing promoter of apoptosis of cancer cell, cancer cell proliferation inhibitor or preventive/therapeutic agent of cancer. (XIII) is useful for producing inhibitor of apoptosis of neuron or preventive/therapeutic agent of neurodegenerative diseases. (I) is useful for screening compounds utilized for promoting or inhibiting apoptosis for treating cancer (e.g. breast or lung cancer) or neurodegenerative diseases (e.g. Alzheimer's disease). (I) enables screening of promoter or inhibitor of apoptosis.

```
XX
SQ
    Sequence 2778 BP; 930 A; 545 C; 605 G; 698 T; 0 U; 0 Other;
                        16.3%; Score 98; DB 15; Length 2778;
 Best Local Similarity
                        100.0%; Pred. No. 1.5e-06;
 Matches
           98: Conservative
                              0; Mismatches
                                                0; Indels
                                                             0; Gaps
                                                                         0;
Qу
           1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
             909 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 850
Db
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qy
             849 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 812
Db
RESULT 3
AED21168/c
ID
    AED21168 standard; DNA; 2980 BP.
XX
AC
    AED21168;
XX
DT
    15-DEC-2005
                (first entry)
XX
DE
    Human Dlg (disks large) DNA sequence.
XX
KW
    secreted frizzled-related protein; tumor; cytostatic; neoplasm;
KW
    drug screening; skin tumor; lymphoma; hematological disease;
KW
    immune disorder; disks large; Dlg; ds; gene.
XX
os
    Homo sapiens.
XX
FH
                    Location/Qualifiers
    Key
FT
    CDS
                    189. .2903
FT
                    /*tag= a
FT
                    /product= "disks large protein"
XX
PN
    WO2005094887-A1.
XX
PD
    13-OCT-2005.
XX
PF
    30-MAR-2005; 2005WO-JP006163.
XX
PR
    31-MAR-2004; 2004JP-00106315.
XX
PΑ
     (DAUC ) DAIICHI PHARM CO LTD.
XX
ΡI
    Akiyama T, Ishidao T, Aiba T;
XX
DR
    WPI; 2005-725415/74.
    P-PSDB; AED21169.
DR
XX
РТ
    Enhancer of expression and/or function of secreted frizzled-related
PT
    protein for inhibiting tumorigenesis, comprises a compound capable of
PT
    increasing expression and/or function of discs large.
XX
```

The present sequence represents Discs large homolog 1 (DLG1) gene.

CC

```
PS
     Disclosure; SEQ ID NO 1; 54pp; Japanese.
XX
CC
     The new invention relates to the finding that tumor formation and the
     transcription of secreted frizzled-related protein (sFRP) gene is lowered
CC
CC
     in a Dlg-knockout mouse. Specifically claimed is an enhancer (I) of the
CC
     expression and/or function of secreted frizzled-related protein (sFRP),
CC
     comprising a compound capable of increasing the expression and/or
CC
     function of Dlg (discs large). Also claimed are a tumorigenesis inhibitor
CC
     (II) containing (I); an agent (III) for preventing or treating tumor,
CC
     containing (I); increasing (M1) the expression and/or function of sFRP,
CC
     comprising inducing the expression and/or function of Dlg; a non-human
CC
     mammal (IV) having one of the deleted Dlg allele; a cell (V) derived from
CC
     a non-human mammal having a deleted Dlg allele; and detecting tumor
CC
     tissue or cells, comprising measuring the expression and/or function of
CC
     Dlg in a sample of the cell or tissue, comparing the expression and/or
CC
     function with a healthy tissue or a cell and detecting whether the
CC
     expression and/or function is reduced. In (I), the compound that
CC
     increases the expression and/or function of Dlg is chosen from a Dlg gene
CC
     or recombinant vector containing the Dlg gene. The sFRP is sFRP2. (M1)
CC
     involves using a Dlg gene or recombinant vector containing the Dlg gene.
CC
     (I) or (M1) are useful for inhibiting tumorigenesis and for preventing or
CC
     treating tumor. (IV) or (V) is useful for identifying a compound capable
CC
     of increasing the expression and/or function of Dlg or sFRP, or
CC
     inhibiting tumorigenesis. (II) and (III) are useful for treating a tumor
CC
     such as skin tumor or lymphoma. (I) effectively induces the expression
CC
     and/or function of sFRP. The present sequence is human disks large DNA
CC
     sequence.
XX
SQ
     Sequence 2980 BP; 949 A; 595 C; 672 G; 764 T; 0 U; 0 Other;
                         16.3%; Score 98; DB 14; Length 2980;
 Best Local Similarity
                         100.0%; Pred. No. 1.5e-06;
 Matches
                                0; Mismatches
           98; Conservative
                                                 0;
                                                     Indels
                                                                           0;
Qу
           1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
              Db
         1097 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 1038
Qу
           61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
              Db
         1037 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 1000
RESULT 4
AEF75023/c
ID
    AEF75023 standard; DNA; 2980 BP.
XX
AC
    AEF75023;
XX
DT
    06-APR-2006 (first entry)
XX
DE
    Human polynucleotide #537.
XX
KW
    Diagnosis; gene regulation; gene expression;
KW
    post traumatic stress disorder; psychiatric disorder; tranquilizer; qene;
KW
XX
```

os Homo sapiens. XX PN WO2006013561-A2. XX PD 09-FEB-2006. XXPF 02-AUG-2005; 2005WO-IL000824. XX PR 02-AUG-2004; 2004US-0592408P. XX PA (YISS) YISSUM RES DEV CO HEBREW UNIV JERUSALEM. PΑ (HADA-) HADASIT MEDICAL RES SERVICES & DEV LTD. XX PΙ Segman R, Shalev A, Goltser T, Friedman N, Shefi N, Kaminski N; XX DR WPI; 2006-145797/15. XX PТ New kit comprising 10 and no more than 574 polynucleotides capable of PTspecifically binding at least one specific polynucleotide sequence, PT useful for determining predisposition of a subject to develop PTSD, or PT for diagnosing PTSD. XX PS Claim 1; SEQ ID NO 537; 157pp; English. XX CC The invention relates to a kit for determining predisposition of a CC subject to developing post-traumatic stress disorder (PTSD) comprising at CC least 10 and no more than 574 polynucleotides, where each of the CC polynucleotides is capable of specifically binding at least one specific CC polynucleotide sequence. The invention also relates to a kit for CC diagnosing PTSD in a subject, agents for the manufacture of the kits CC cited comprising the polynucleotides cited, and a microarray comprising CC at least 10 and no more than 904 oligonucleotides where each of the CC oligonucleotides is capable of specifically binding at least one specific CC polynucleotide sequence. The kit comprises each of the polynucleotides CC selected from an oligonucleotide molecule, a cDNA molecule, a genomic CC molecule and an RNA molecule. Each of the polynucleotides is at least 10 CC and no more than 50 nucleic acids in length. Each of the polynucleotides CC is bound to a solid support. The kit also comprises at least one reagent CC suitable for detecting hybridization of the polynucleotides and at least CC one RNA transcript. The kit further comprises packaging materials CC packaging the at least one reagent and instructions for using the kit in CC determining predisposition of the subject to developing PTSD, or for CC diagnosing the disease. The microarray comprises oligonucleotides of at CC least 10 and no more than 40 nucleic acids in length. The agent is CC capable of regulating an expression level of at least one gene as a CC pharmaceutical or for the manufacture of a medicament identified for CC preventing PTSD. The kit is useful for determining predisposition of a CC subject to developing PTSD or for diagnosing PTSD. This sequence CC represents a human polynucleotide of the invention. Note: The sequence CC data for this patent did not form part of the printed specification, but CC was obtained in electronic format directly from WIPO at CC ftp.wipo.int/pub/published pct sequences. XX SO Sequence 2980 BP; 949 A; 595 C; 672 G; 764 T; 0 U; 0 Other;

Query Match 16.3%; Score 98; DB 15; Length 2980; Best Local Similarity 100.0%; Pred. No. 1.5e-06;

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Matches
           98; Conservative
                               0; Mismatches
                                                0; Indels
                                                              0; Gaps
                                                                         0;
Qу
           1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
             Db
        1097 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 1038
Qу
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
             Db
        1037 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 1000
.....
RESULT 5
ACC85071/c
    ACC85071 standard; DNA; 3046 BP.
XX
AC
    ACC85071;
XX
DT
    13-OCT-2003 (first entry)
XX
DΕ
    Human MBCAT polypeptide encoding DNA.
XX
KW
    MBCAT; beta-catenin; cytostatic; gene therapy; cancer; human; gene; ds.
XX
os
    Homo sapiens.
XX
FH
                    Location/Qualifiers
    Key
FT
    CDS
                    189. .2969
FT
                    /*tag= a
                    /product= "MBCAT"
FT
XX
PN
    WO2003052068-A2.
XX
PD
    26-JUN-2003.
XX
PF
    12-DEC-2002; 2002WO-US039796.
XX
PR
    13-DEC-2001; 2001US-0340213P.
PR
    13-DEC-2001; 2001US-0340314P.
PR
    13-DEC-2001; 2001US-0340322P.
PR
    15-FEB-2002; 2002US-0357502P.
XX
PA
     (EXEL-) EXELIXIS INC.
XX
PΙ
    Costa MA, Gendreau SB, Dora EG, Nicoll M;
XX
DR
    WPI; 2003-533010/50.
DR
    P-PSDB; ABR82220.
XX
PT
     Identifying a candidate beta-catenin pathway modulating agent for
PT
     diagnosing or treating cancer by detecting a test agent-biased activity
PT
    of the assay system comprising a purified MBCAT polypeptide or nucleic
PT
    acid.
XX
PS
    Example; Page 49-51; 81pp; English.
XX
CC
     The invention relates to genes that modify beta-catenin pathway and to
CC
     the identification of human MBCAT (modifiers of beta-catenin) genes. The
```

```
CC
     beta-catenin function and are useful for manufacturing a medicament for
CC
     diagnosing or treating breast, colon, lung or ovary cancer. The present
CC
     sequence represents a human MBCAT polypeptide encoding DNA
XX
SQ
     Sequence 3046 BP; 975 A; 612 C; 681 G; 778 T; 0 U; 0 Other;
  Query Match
                         16.3%; Score 98; DB 9; Length 3046;
                         100.0%; Pred. No. 1.5e-06;
  Best Local Similarity
  Matches
           98; Conservative
                               0; Mismatches
                                                 0;
                                                     Indels
                                                              0;
                                                                          0;
                                                                  Gaps
           1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qу
             Db
         1097 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 1038
           61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qy
             1037 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 1000
Db
RESULT 6
ADK11476/c
     ADK11476 standard; DNA; 3046 BP.
XX
AC
    ADK11476;
XX
DT
     06-MAY-2004 (first entry)
XX
DE
     Human discs large (Drosophila) homolog 1 gene.
XX
KW
     ds; gene; cytostatic; cardiovascular; immunosuppressive; nephrotropic;
KW
     antirheumatic; antiarthritic; dermatological; antipsoriatic;
KW
     antiinflammatory; fungicide; gene therapy; Drosophila; diagnosis;
KW
     cardiovascular disorder; autoimmune disease; glomerulonephritis;
KW
     rheumatoid arthritis; dermatological disorder; psoriasis;
KW
     inflammatory disorder; malaria; emphysema; alopecia.
XX
os
     Homo sapiens.
XX
PN
     WO2003040301-A2.
XX
PD
     15-MAY-2003.
XX
PF
     23-OCT-2002; 2002WO-GB004780.
XX
PR
     05-NOV-2001; 2001GB-00026506.
PR
     27-NOV-2001; 2001GB-00028384.
PR
     11-FEB-2002; 2002GB-00003185.
XX
PA
     (CYCL-) CYCLACEL LTD.
XX
ΡI
     Deak P, Frenz L, Glover D, Midgley C;
XX
DR
     WPI; 2003-441540/41.
DR
     P-PSDB; ADK11477.
XX
PT
     New Drosophila polypeptides and polynucleotides, useful for diagnosing,
```

MBCAT polypeptides are therapeutic targets for disorders associated with

CC

```
PT
     preventing and/or treating disorders, such as cancer, glomerulonephritis,
PT
     rheumatoid arthritis, psoriasis, malaria, emphysema and alopecia.
XX
PS
     Example 28; Page 225; 265pp; English.
XX
CC
     The invention relates to novel Drosophila species DNA sequences and their
CC
     encoded proteins with their corresponding human homologues. The proteins
CC
     or their encoding polynucleotides are useful in a method of prevention,
CC
     treatment or diagnosis of a disease in an individual, and used to
CC
     identify a substance capable of binding to the polypeptide or modulating
CC
     the function of the polypeptide comprising incubating the polypeptide
     with a candidate substance and determining whether the substance binds to
CC
CC
     the polypeptide. The compositions are administered to an individual in
     need of such treatment. The method of diagnosis, in which the presence or
CC
     absence of a polynucleotide is detected in a biological sample, comprises
CC
CC
     brining the biological sample containing the nucleic acid such as DNA or
CC
     RNA into contact with a probe comprising a fragment of at least 15
CC
     nucleotides of the polynucleotide, and detecting any duplex formed
CC
     between the probe and nucleic acid in the sample. The method also
CC
     comprises providing an antibody capable of binding to the polypeptide,
CC
     incubating a biological sample with the antibody to allow the formation
CC
     of an antibody-antigen complex, and determining whether antibody-antigen
CC
     complex comprising the antibody is formed. The disease comprises a
CC
     proliferative disease such as cancer. The antibody or identified
     substance is also useful in inhibiting the function of a polypeptide
CC
CC
     and/or regulating a cell division cycle function. The diseases also
CC
     include cardiovascular disorders, autoimmune diseases such as
     glomerulonephritis and rheumatoid arthritis, and dermatological disorders
CC
     such as psoriasis, inflammatory, fungal, and parasitic disorders such as
CC
CC
     malaria, emphysema and alopecia. This sequence represents a human homolog
CC
     gene for one of the Drosophila genes of the invention.
XX
SQ
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  Query Match
                         16.3%; Score 98; DB 10; Length 3046;
  Best Local Similarity
                         100.0%; Pred. No. 1.5e-06;
  Matches
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                                0; Mismatches
                                                  0:
                                                     Indels
                                                               0;
                                                                   Gaps
                                                                           0;
Qy
           1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTTTACATCACGAAC 60
              Db
         1097 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 1038
Qу
           61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
              Db
         1037 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 1000
RESULT 7
ADR83538/c
ID
     ADR83538 standard; DNA; 3046 BP.
XX
AC
    ADR83538;
XX
DT
     02-DEC-2004
                (first entry)
XX
DE
    Human discs large homolog 1 DNA, target gene of miRNA.
XX
```

```
KW
     human; ds; miRNA; microRNA; ontogenesis; cell therapy; cancer;
KW
     immune disease; nerve disorder; amyotrophic lateral sclerosis;
KW
     Parkinson's disease; Alzheimer's disease; inflammatory disease;
KW
     siRNA silencing precursor; cytostatic; immunosuppressive; nootropic;
KW
     neuroprotective; antiinflammatory; immunotherapy; discs large homolog 1.
XX
os
     Homo sapiens.
XX
PN
     WO2004076622-A2.
XX
PD
     10-SEP-2004.
XX
PF
     10-FEB-2004; 2004WO-JP001433.
XX
PR
     10-FEB-2003; 2003US-0445829P.
XX
PA
     (NAAD-) NAT INST ADVANCED IND SCI & TECHNOLOGY.
XX
PΙ
     Taira K, Kawasaki H;
XX
DR
     WPI; 2004-653393/63.
XX
PT
     Modulating expression of a target gene in a cell, for treating cancer, an
PT
     immune disease, or a nerve disorder, comprises introducing into the cell
PT
     a polynucleotide that forms a duplex region with an mRNA transcribed from
PT
     the target gene.
XX
PS
     Claim 9; SEQ ID NO 440; 865pp; English.
XX
CC
     This invention relates to a novel method for modulating the expression of
CC
     a target gene in a cell. Specifically, it refers to the introduction into
CC
     a cell of a polynucleotide that forms a duplex region with an mRNA
CC
     transcribed from the target gene, where the duplex region comprises a
CC
     mammalian miRNA target region i.e. a non-coding microRNA (miRNA) that
CC
     regulates mRNA at a post-transcriptional level. The present invention
CC
     describes a method for controlling ontogenesis of a mammal, function of a
CC
     mammalian cell, differentiation of a mammalian cell or viability of a
CC
     mammalian cell in the post-transcriptional phase, which comprises
CC
     introducing a plasmid vector comprising a promoter and nucleic acid
CC
     molecule expressing an miRNA or siRNA silencing precursor to the miRNA.
CC
     Accordingly, it provides a cell therapy method for treating cancer,
CC
     immune disease, nerve disorder (e.g. amyotrophic lateral sclerosis,
CC
     Parkinson's disease, or Alzheimer's disease) or an inflammatory disease
CC
     by introducing into the cell the miRNA, siRNA silencing precursor to the
CC
     miRNA or the plasmid vector. As such, they can be developed into
CC
     pharmaceutical compositions that exhibit cytostatic, immunosuppressive,
CC
     nootropic, neuroprotective and antiinflammatory activities and hence can
CC
     be used for immunotherapy. This polynucleotide sequence is a human target
CC
     gene whose expression is modulated by miRNAs of the invention.
XX
SQ
     Sequence 3046 BP; 975 A; 612 C; 681 G; 778 T; 0 U; 0 Other;
  Query Match
                          16.3%; Score 98; DB 13; Length 3046;
  Best Local Similarity
                          100.0%; Pred. No. 1.5e-06;
 Matches
            98; Conservative
                                 0; Mismatches
                                                       Indels
```

Qу

```
1097 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 1038
Db
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qу
             Db
        1037 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 1000
RESULT 8
ADU05806/c
ID
    ADU05806 standard; DNA; 3046 BP.
XX
AC
    ADU05806;
XX
DT
    27-JAN-2005 (first entry)
XX
DE
    Novel bronchial cancer-associated human gene SeqID28.
XX
KW
    bronchial cancer; cytostatic; tumour-associated protein;
KW
    cancer detection; metastasis; tumour; gene; ds; human.
XX
os
    Homo sapiens.
XX
PN
    DE10316701-A1.
XX
PD
    04-NOV-2004.
XX
ΡF
    09-APR-2003; 2003DE-01016701.
XX
PR
    09-APR-2003; 2003DE-01016701.
XX
PA
     (HINZ/) HINZMANN B.
     (HERM/) HERMANN K.
PA
     (CAST/) HEIDEN CASTANOS-VELEZ E.
PA
XX
    Mennerich D, Bruemmendorf T, Heiden E, Hermann K, Kinnemann H;
PΙ
PΙ
    Li X, Roepcke S, Staub E, Hinzmann B, Rosenthal A, Pilarsky C;
XX
DR
    WPI; 2004-786403/78.
DR
    P-PSDB; ADU06293.
XX
PT
    New nucleic acid, and derived proteins, useful for diagnosis of bronchial
PT
    cancer and in screening for therapeutic and diagnostic agents.
XX
PS
    Claim 1; SEQ ID NO 28; 1381pp; German.
XX
CC
    This invention relates to a novel isolated nucleic acid associated with
CC
    bronchial cancer comprising 489 defined sequences given in the
CC
     specification. The invention may be useful for the production of
CC
    compounds with a cytostatic activity through the inhibition of expression
CC
    or activity of tumour-associated proteins. The novel DNA sequences and
CC
    the proteins/peptides encoded by them are used for detecting bronchial
CC
    cancer or determining the risk of developing it and to screen for
CC
    specific binding partners of the DNA or protein sequences, where the
CC
    binding partners are potentially useful as agents for treating or
    diagnosing bronchial cancer. The DNA or protein sequences can also be
CC
CC
    used for prognosis, detection of metastases and for secondary treatment
```

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OM nucleic - nucleic search, using sw model

Run on: July 15, 2006, 18:16:32; Search time 1201 Seconds

(without alignments)

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Searched: 18892170 seqs, 6143817638 residues

Total number of hits satisfying chosen parameters: 37784340

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Maximum Match 100%

Listing first 100 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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ALIGNMENTS

RESULT 1

US-10-722-939-1

- ; Sequence 1, Application US/10722939
- ; Publication No. US20050214771A1
- ; GENERAL INFORMATION:
- ; APPLICANT: ROTH, RICHARD B.
- ; APPLICANT: NELSON, MATTHEW ROBERTS
- ; APPLICANT: KAMMERER, STEFAN M.
- ; APPLICANT: BRAUN, ANDREAS
- ; APPLICANT: RENELAND, RIKARD

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TITLE OF INVENTION: METHODS FOR IDENTIFYING RISK OF BREAST CANCER AND
TREATMENTS
  TITLE OF INVENTION: THEREOF
  FILE REFERENCE: SEQ-4071-UT
  CURRENT APPLICATION NUMBER: US/10/722,939
  CURRENT FILING DATE: 2003-11-26
  PRIOR APPLICATION NUMBER: 60/429,136
  PRIOR FILING DATE: 2002-11-25
  PRIOR APPLICATION NUMBER: 60/490,234
  PRIOR FILING DATE: 2003-07-23
  NUMBER OF SEQ ID NOS: 634
  SOFTWARE: PatentIn version 3.2
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  TYPE: DNA
  ORGANISM: Homo sapiens
US-10-722-939-1
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; Publication No. US20040181048A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single
  TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.135
  CURRENT APPLICATION NUMBER: US/09/925,065A
  CURRENT FILING DATE: 2001-08-08
  PRIOR APPLICATION NUMBER: US 60/243,096
  PRIOR FILING DATE: 2000-10-24
  PRIOR APPLICATION NUMBER: US 60/252,147
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/250,092
  PRIOR FILING DATE: 2000-11-30
  PRIOR APPLICATION NUMBER: US 60/261,766
  PRIOR FILING DATE: 2001-01-16
  PRIOR APPLICATION NUMBER: US 60/289,846
  PRIOR FILING DATE: 2001-05-09
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; Publication No. US20050228172A9
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single
  TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.135
  CURRENT APPLICATION NUMBER: US/09/925,065A
  CURRENT FILING DATE: 2001-08-08
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  PRIOR FILING DATE: 2000-10-24
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- ; Publication No. US20050227243A1
- ; GENERAL INFORMATION:
- ; APPLICANT: Cyclacel Limited
- ; APPLICANT: Deak, Peter
- ; APPLICANT: Frenz, Lisa
- ; APPLICANT: Glover, David
- ; APPLICANT: Midgley, Carol
- ; TITLE OF INVENTION: Cell Cycle Progression Proteins
- ; FILE REFERENCE: 10069/2012
- CURRENT APPLICATION NUMBER: US/10/840,060
- ; CURRENT FILING DATE: 2004-05-05
- ; PRIOR APPLICATION NUMBER: PCT/GB02/04780
- PRIOR FILING DATE: 2002-10-23
- ; PRIOR APPLICATION NUMBER: GB 0126506.5

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  PRIOR FILING DATE: 2001-11-27
  PRIOR APPLICATION NUMBER: GB 0203185.4
  PRIOR FILING DATE: 2002-02-11
  NUMBER OF SEQ ID NOS: 306
  SOFTWARE: PatentIn version 3.1
 SEO ID NO 266
   LENGTH: 3046
   TYPE: DNA
   ORGANISM: Homo sapiens
US-10-840-060-266
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 Best Local Similarity 100.0%; Pred. No. 1e-06;
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; Sequence 5, Application US/10722939
; Publication No. US20050214771A1
; GENERAL INFORMATION:
  APPLICANT: ROTH, RICHARD B.
  APPLICANT: NELSON, MATTHEW ROBERTS
  APPLICANT: KAMMERER, STEFAN M.
  APPLICANT: BRAUN, ANDREAS
  APPLICANT: RENELAND, RIKARD
  TITLE OF INVENTION: METHODS FOR IDENTIFYING RISK OF BREAST CANCER AND
TREATMENTS
; TITLE OF INVENTION: THEREOF
; FILE REFERENCE: SEQ-4071-UT
  CURRENT APPLICATION NUMBER: US/10/722,939
  CURRENT FILING DATE: 2003-11-26
  PRIOR APPLICATION NUMBER: 60/429,136
  PRIOR FILING DATE: 2002-11-25
  PRIOR APPLICATION NUMBER: 60/490,234
  PRIOR FILING DATE: 2003-07-23
  NUMBER OF SEQ ID NOS: 634
  SOFTWARE: PatentIn version 3.2
; SEQ ID NO 5
   LENGTH: 3046
   TYPE: DNA
   ORGANISM: Homo sapiens
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PRIOR FILING DATE: 2001-11-05

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             Db
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; Sequence 1545, Application US/10106698
; Publication No. US20030109690A1
; GENERAL INFORMATION:
  APPLICANT: Ruben et al.
  TITLE OF INVENTION: Colon and Colon Cancer Associated Polynucleotides and
Polypeptides
  FILE REFERENCE: PA005P1
  CURRENT APPLICATION NUMBER: US/10/106,698
  CURRENT FILING DATE: 2002-03-27
  PRIOR APPLICATION NUMBER: PCT/US00/26524
  PRIOR FILING DATE: 2000-09-28
  PRIOR APPLICATION NUMBER: US 60/157,137
  PRIOR FILING DATE: 1999-09-29
  PRIOR APPLICATION NUMBER: US 60/163,280
  PRIOR FILING DATE: 1999-11-03
  NUMBER OF SEQ ID NOS: 8564
  SOFTWARE: PatentIn Ver. 3.0
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   ORGANISM: Homo sapiens
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; Sequence 11234, Application US/09960352
; Patent No. US20020137139A1
; GENERAL INFORMATION:
  APPLICANT: Warren, Wesley C.
  APPLICANT: Tao, Nengbing
  APPLICANT: Byatt, John C.
  APPLICANT: Mathialagan, Nagappan
  TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH
LACTATION AND
  TITLE OF INVENTION: MUSCLE AND FAT DEPOSITION
 FILE REFERENCE: 16511.006/37-21(10298)C
  CURRENT APPLICATION NUMBER: US/09/960,352
  CURRENT FILING DATE: 2001-09-24
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  LENGTH: 419
  TYPE: DNA
  ORGANISM: Bos taurus
  OTHER INFORMATION: Clone ID: 48-LIB3058-052-Q1-K1-D8
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RESULT 8 US-10-473-126-386/c

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; Publication No. US20040234973A1
GENERAL INFORMATION:
 APPLICANT: Epigenomics AG
 TITLE OF INVENTION: Methods and nucleic acids for the analysis of
hematopoietic cell
 TITLE OF INVENTION: proliferative disorders
 FILE REFERENCE:
 CURRENT APPLICATION NUMBER: US/10/473,126
 CURRENT FILING DATE: 2003-09-26
 NUMBER OF SEQ ID NOS: 1258
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  LENGTH: 8056
  TYPE: DNA
  ORGANISM: Artificial Sequence
  FEATURE:
  OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
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; Sequence 7873, Application US/10425115
; Publication No. US20040214272A1
; GENERAL INFORMATION:
 APPLICANT: La Rosa, Thomas J.
 APPLICANT: Kovalic, David K.
 APPLICANT: Zhou, Yihua
 APPLICANT: Cao, Yongwei
 TITLE OF INVENTION: Nucleic Acid Molecules and Other Molecules Associated
With
 TITLE OF INVENTION: Plants
; FILE REFERENCE: 38-21(53222)B
 CURRENT APPLICATION NUMBER: US/10/425,115
 CURRENT FILING DATE: 2003-04-28
; NUMBER OF SEQ ID NOS: 369326
; SEQ ID NO 7873
  LENGTH: 542
  TYPE: DNA
  ORGANISM: Zea mays
  FEATURE:
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US-10-425-115-7873
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; Sequence 306, Application US/10486706
; Publication No. US20050071088A1
; GENERAL INFORMATION:
  APPLICANT: LANDFIELD, PHILIP W.
  APPLICANT: BLALOCK, ERIC M.
  APPLICANT: CHEN, KUEY-CHU
  APPLICANT: FOSTER, THOMAS C.
  TITLE OF INVENTION: GENE EXPRESSION PROFILE BIOMARKERS AND THERAPEUTIC
TARGETS FOR
  TITLE OF INVENTION: BRAIN AGING AND AGE-RELATED COGNITIVE IMPAIRMENT
  FILE REFERENCE: 50229-426
  CURRENT APPLICATION NUMBER: US/10/486,706
  CURRENT FILING DATE: 2004-02-13
  PRIOR APPLICATION NUMBER: PCT/US02/25607
  PRIOR FILING DATE: 2002-08-13
  PRIOR APPLICATION NUMBER: US 60/311,343
  PRIOR FILING DATE: 2001-08-13
  NUMBER OF SEQ ID NOS: 461
  SOFTWARE: PatentIn version 3.2
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   LENGTH: 3256
   TYPE: DNA
   ORGANISM: Rattus norvegicus
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; Sequence 14, Application US/10980571
; Publication No. US20050138675A1
; GENERAL INFORMATION:
  APPLICANT: Pfizer, Inc.
  TITLE OF INVENTION: Method For Determining Cardiotoxicity
  FILE REFERENCE: PC26200A
  CURRENT APPLICATION NUMBER: US/10/980,571
  CURRENT FILING DATE: 2004-11-02
  PRIOR APPLICATION NUMBER: 60/516,774
  PRIOR FILING DATE: 2003-11-03
  NUMBER OF SEQ ID NOS: 18
  SOFTWARE: PatentIn version 3.3
; SEQ ID NO 14
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   TYPE: DNA
   ORGANISM: Rattus norvegicus
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; Sequence 240, Application US/10473126
; Publication No. US20040234973A1
; GENERAL INFORMATION:
  APPLICANT: Epigenomics AG
  TITLE OF INVENTION: Methods and nucleic acids for the analysis of
hematopoietic cell
  TITLE OF INVENTION: proliferative disorders
  FILE REFERENCE:
  CURRENT APPLICATION NUMBER: US/10/473,126
  CURRENT FILING DATE: 2003-09-26
  NUMBER OF SEQ ID NOS: 1258
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   LENGTH: 8056
   TYPE: DNA
  ORGANISM: Artificial Sequence
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
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 Best Local Similarity
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Searched: 48236798 seqs, 27959665780 residues

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Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 100 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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С	24	82.8	13.8	867	14	CNS00CX5	AL060052 Drosophil
	25	82.6	13.8	866	5	CF289423	CF289423 AGENCOURT
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	38	78.6	13.1	1101	14	CNS003BD	AL064091 Drosophil
	39	78.4	13.1	759	14	CNS06QXV	AL411257 T7 end of
С	40	78.4	13.1	1007	10	DV050335	DV050335 DAY35S_08
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C	46	77.6	12.9	1283	13		CL641341 CH213-901
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	76	74.8	12.5	1522	12	CL128484	CL128484 ISB1-94I1
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	80	74.6	12.4	1058	12	CL077132	CL077132 CH216-143
	81	74.6	12.4	1101	14	CNS0039G	AL063921 Drosophil
С	82	74.6	12.4	1542	14	AG386981	AG386981 Mus muscu
	83	74.4	12.4	734	14	CNS010MP	AL099163 Drosophil
С	84	74.4	12.4	749	11	AQ324504	AQ324504 mgxb0018L
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	86	74.4	12.4	945	14	CNS04D0K	AL285149 Tetraodon
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С	88	74.4	12.4	1027	14	CNS02T50	AL212733 Tetraodon
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ALIGNMENTS

KEYWORDS EST.
SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

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Hominidae; Homo.
REFERENCE
             (bases 1 to 740)
 AUTHORS
          NIH-MGC http://mgc.nci.nih.gov/.
          National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
 JOURNAL
          Unpublished (1999)
COMMENT
          Contact: Robert Strausberg, Ph.D.
          Email: cgapbs-r@mail.nih.gov
          Tissue Procurement: CLONETECH Laboratories, Inc.
           cDNA Library Preparation: CLONETECH Laboratories, Inc.
           cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
           DNA Sequencing by: Incyte Genomics, Inc.
           Clone distribution: MGC clone distribution information can be
          found through the I.M.A.G.E. Consortium/LLNL at:
          http://image.llnl.gov
          Plate: LLCM1561 row: a column: 09
          High quality sequence stop: 474.
FEATURES
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                  /organism="Homo sapiens"
                  /mol type="mRNA"
                  /db xref="taxon:9606"
                  /clone="IMAGE:4715024"
                  /lab host="DH10B (T1 phage-resistant)"
                  /clone lib="NIH MGC 76"
                  /note="Organ: liver; Vector: pDNR-LIB (Clontech); Site 1:
                  SfiI (ggccgcctcggcc); Site 2: SfiI (ggccattatggcc); 5' and
                  3' adaptors were used in cloning as follows: 5' adaptor
                  sequence: 5'-CACGGCCATTATGGCC-3' and 3' adaptor sequence:
                  5'-ATTCTAGAGGCCGAGGCGGCCGACATG-dT(30)BN-3' (where B = A,
                  C, or G and N = A, C, G, or T). Average insert size 1.85
                  kb (range 1.0-4.0 kb). 15/15 colonies contained inserts
                  by PCR. This library was enriched for full-length clones
                  and was constructed by Clontech Laboratories (Palo Alto,
                  CA). Note: this is a NIH MGC Library."
ORIGIN
                             Score 138.4; DB 2; Length 740;
 Query Match
                       23.1%;
 Best Local Similarity
                       99.3%;
                             Pred. No. 8.6e-14;
 Matches 139; Conservative
                             0; Mismatches
                                                Indels
                                                                   0:
Qу
        Db
        521 TCACAGATGGAGAAACTGAGGCACAAAGAATGTAAATAACTTTCCTAAGGCCACCCAGAT 580
Qу
            Db
        392 TCACAGATGGAGAAACTGAGGCACAAAGAATGTAAATAACTTTCCTAAGGCCACCCAGAT 333
        581 AATAAGTGACAGAGCTGTGA 600
Qy
            Db
        332 AATAAGTGACAGAGCTGTGA 313
RESULT 2
CC504900
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LOCUS CC504900 906 bp DNA linear GSS 17-JUN-2003 DEFINITION CH240_345A2.T7 CHORI-240 Bos taurus genomic clone CH240_345A2,

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genomic survey sequence.
            CC504900
ACCESSION
VERSION
            CC504900.1 GI:31823193
KEYWORDS
            GSS.
SOURCE
            Bos taurus (cattle)
  ORGANISM Bos taurus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Laurasiatheria; Cetartiodactyla; Ruminantia;
            Pecora; Bovidae; Bovinae; Bos.
REFERENCE
            1 (bases 1 to 906)
  AUTHORS
            Holt, R., Stott, J., Yang, G., Barber, S., Smailus, D., Prabhu, A.-L.,
            Tsai, M., Cloutier, A., Lee, D., Girn, N., Olson, T., Mayo, M.,
            Butterfield, Y., Kirkpatrick, R., Liu, J., Guin, R., Chan, A., Chiu, R.,
            Mathewson, C., Wye, N., Masson, A., Brown-John, M., Jones, S.,
            Schein, J., Marra, M., de Jong, P., McWilliam, S., Barris, W.,
            Dalrymple, B.P. and Tellam, R.
  TITLE
            Bovine BAC End Sequences from Library CHORI-240, PLATES 294 to 398
  JOURNAL
            Unpublished (2003)
COMMENT
            Other GSSs: CH240 345A2.TARBAC13P2
            Contact: Rob Holt
            Sequencing
            The British Columbia Cancer Agency Genome Science Centre
            600 W. 10th Ave, Vancouver, British Columbia, Canada V5Z 4E6
            Tel: 604-877-6085
            Fax: 604-877-6276
            Email: rholt@bcgsc.ca
            Clones are derived from the bovine BAC library CHORI-240
            (http://www.chori.org/bacpac/bovine240.htm). For BAC library
            availability, please contact Pieter de Jong (pdejong@mail.cho.org).
            Clones may be purchased from BACPAC Resources
            (http://www.chori.org/bacpac/ordering information.htm). This work
            was undertaken as part of the International Bovine BAC Mapping
            Consortium (IBBMC) by CSIRO Livestock Industries, Australia and the
            British Columbia Genome Sciences Centre, Canada.
            Plate: 345 row: A column: 2
            Seq primer: T7
            Class: BAC ends.
FEATURES
                     Location/Qualifiers
     source
                     1. .906
                     /organism="Bos taurus"
                     /mol type="genomic DNA"
                     /strain="breed: Hereford"
                     /db xref="taxon:9913"
                     /clone="CH240 345A2"
                     /sex="Male"
                     /cell type="Blood"
                     /clone lib="CHORI-240"
                     /note="Vector: pTARBAC1.3; Site_1: MboI; Site_2: MboI;
                     Hereford bull L1 Domino 99375; CHORI-240 Bovine BAC
                     library (Male) produced by Pieter de Jong"
ORIGIN
  Query Match
                          22.8%; Score 136.8; DB 12; Length 906;
  Best Local Similarity
                          63.6%; Pred. No. 1.6e-13;
 Matches 347; Conservative
                                 0; Mismatches 147; Indels
                                                                 52;
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1 ACGAACATCTATTTCATTTACTCGCAAAATACAGTCATTGACTCTAAGGAAA-AATTAAA 59
Db
        Qу
            60 AATAAAAATGAAATATTAA----AAATTTAATATCGATTTTTCACTTTGCCAATTTTTAT 115
Db
        175 TTACTTCTTTTTAAGGTAAAGAGAATTATAAATAATTCTGGAGTAATTCCAGAAAACAT 234
Qу
                Db
        116 TTTCACATTTTTAAGGTTAACTACTACGATTTTTTCTGAGAAGAAACTTCATGTTAA--- 172
        235 AAATGAAGAAGTATATCAAAAACTAATATAAACAAATACAAACATTTCCCAAGGGCCAG 294
Qу
                  Db
        173 ----ATTTCAATATATCAAATACTTATAT----AAATACAAACATTTCTCAAGGGCCAT 223
        Qу
           Db
        224 CAAAGGAAATAGTGCAAAT-----AATAGATTATTTTACAAAAATATTAAATGATAATG 277
        355 ACAGCTACCATTTGCATACTTCCTGCATGCTTATTCTGTGTAGGCATTAAGACACACTTT 414
Qу
           278 ACAGCT-----
                                -----GTCTATGTACAGAAATTAAAATACTTT 310
Db
        415 ATAAAAATAGCAAACATTTATTTAGCACTAACCACATGCCAGGCACTTTCTTGGTATTTT 474
Qу
           311 AGAAAAACAGAAAATACTTATTTAGTACTAATCACAAGCTACAAACTTCTTTTAGTGTTT 370
Db
        475 AACCCTCATGACACCTGTAAGCTTAATATATATTTTAATCCCTATTTCACAGATGGAGAA 534
Qу
                                11 111 1 1111111111 111 111 1111 11
                  Db
        371 TAACCCTTACAACAACCTGTAAGGTGGATCTATCTATATCCCTATTTTACACATGGAAAA 430
        535 ACTGAGGCACAAAGAATGTAAATAACTTTCCTAAGGCCACCCAGATAATAAGTGACAGAG 594
Qу
           431 ACTGAGGCACAAAG--TTTAAATAACATTCCCAAAGCCACCCAGATAATGACCAGTAGAG 488
Db
        595 CTGTGA 600
Qу
           | | | | | |
Db
        489 CAGTGA 494
RESULT 3
H96495
LOCUS
         H96495
                             462 bp
                                     mRNA
                                            linear
                                                   EST 25-NOV-1996
DEFINITION
         yt98h02.rl Soares_pineal_gland_N3HPG Homo sapiens cDNA clone
         IMAGE:232371 5' similar to SP:A45436 A45436
         SAP90=SYNAPSE-ASSOCIATED PROTEIN - ;, mRNA sequence.
ACCESSION
         H96495
VERSION
         H96495.1 GI:1109963
KEYWORDS
         EST.
SOURCE
         Homo sapiens (human)
 ORGANISM
         Homo sapiens
         Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
         Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
         Hominidae; Homo.
REFERENCE
            (bases 1 to 462)
 AUTHORS
         Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M.,
         Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M.,
```

```
Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F.,
           Trevaskis, E., Waterston, R., Williamson, A., Wohldmann, P. and
           Wilson, R.
  TITLE
           The WashU-Merck EST Project
  JOURNAL
           Unpublished (1995)
COMMENT
           Contact: Wilson RK
           Washington University School of Medicine
           4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
           Tel: 314 286 1800
           Fax: 314 286 1810
           Email: est@watson.wustl.edu
           High quality sequence stops: 333
           Source: IMAGE Consortium, LLNL
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           Possible reversed clone: similarity on wrong strand
           Insert Length: 2241
                               Std Error: 0.00
           Seq primer: M13RP1.
FEATURES
                   Location/Qualifiers
    source
                   1. .462
                   /organism="Homo sapiens"
                    /mol type="mRNA"
                   /db xref="GDB:3862362"
                   /db_xref="taxon:9606"
                    /clone="IMAGE:232371"
                   /lab host="DH10B (ampicillin resistant)"
                   /clone lib="Soares_pineal_gland_N3HPG"
                   /note="Organ: pineal gland; Vector: pT7T3D (Pharmacia)
                   with a modified polylinker; Site_1: Not I; Site_2: Eco RI;
                   1st strand cDNA was primed with a Not I - oligo(dT) primer
                    3'], double-stranded cDNA was size selected, ligated to
                   Eco RI adapters (Pharmacia), digested with Not I and
                   cloned into the Not I and Eco RI sites of a modified pT7T3
                   vector (Pharmacia). Library constructed by Bento Soares
                   and M.Fatima Bonaldo. "
ORIGIN
 Query Match
                        16.1%; Score 96.8; DB 10;
                                                   Length 462;
 Best Local Similarity
                        98.0%; Pred. No. 1.1e-06;
 Matches
           98; Conservative
                               0; Mismatches
                                                2;
                                                   Indels
                                                             0;
                                                                 Gaps
                                                                        0;
Qу
           1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
             Db
         199 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 258
Qу
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCCTG 100
             Db
         259 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCCGG 298
RESULT 4
AA281170
LOCUS
           AA281170
                                   361 bp
                                            mRNA
                                                   linear
                                                            EST 14-AUG-1997
DEFINITION
           zt01b11.rl NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:711837 5'
           similar to TR:G558438 G558438 HOMOLOG OF DROSOPHILA DISCS LARGE
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PROTEIN, ISOFORM 1. ;, mRNA sequence.

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AA281170.1 GI:1923851
VERSION
KEYWORDS
           EST.
           Homo sapiens (human)
SOURCE
 ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae: Homo.
              (bases 1 to 361)
REFERENCE
 AUTHORS
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
 JOURNAL
           Unpublished (1997)
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           This clone is available royalty-free through LLNL; contact the
           IMAGE Consortium (info@image.llnl.gov) for further information.
           Possible reversed clone: similarity on wrong strand
           Insert Length: 1961
                               Std Error: 0.00
           Seg primer: -28m13 rev2 ET from Amersham
           High quality sequence stop: 360.
FEATURES
                   Location/Qualifiers
                   1. .361
    source
                   /organism="Homo sapiens"
                   /mol_type="mRNA"
                   /db xref="taxon:9606"
                   /clone="IMAGE:711837"
                   /tissue_type="germinal center B cell"
                   /lab_host="DH10B"
                   /clone lib="NCI CGAP GCB1"
                   /note="Vector: pT7T3D-PacI; Site_1: Not I; Site_2: Eco RI;
                   1st strand cDNA was prepared from human tonsillar cells
                   enriched for germinal center B cells by flow sorting
                   (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr.
                   David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA
                   synthesis was primed with a Not I - oligo(dT) primer
                    ]. Double-stranded cDNA was ligated to Eco RI adaptors
                   (Pharmacia), digested with Not I and cloned into the Not I
                   and Eco RI sites of the modified pT7T3 vector. Library
                   went through one round of normalization, and was
                   constructed by Bento Soares and M. Fatima Bonaldo."
ORIGIN
 Query Match
                        16.1%;
                               Score 96.4; DB 1; Length 361;
 Best Local Similarity
                        99.0%; Pred. No. 1.3e-06;
           97; Conservative
                               0; Mismatches
                                               1;
                                                   Indels
                                                                Gaps
                                                                        0;
           1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qy
             Db
          34 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 93
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qy
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94 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 131

ACCESSION

Db

AA281170

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DA501101/c
LOCUS
            DA501101
                                     362 bp
                                               mRNA
                                                       linear
                                                                EST 08-NOV-2005
           DA501101 FCBBF3 Homo sapiens cDNA clone FCBBF3015237 5', mRNA
DEFINITION
            sequence.
ACCESSION
            DA501101
VERSION
            DA501101.1 GI:81184123
KEYWORDS
SOURCE
            Homo sapiens (human)
  ORGANISM
           Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
               (bases 1 to 362)
            Kimura, K., Wakamatsu, A., Suzuki, Y., Ota, T., Nishikawa, T.,
  AUTHORS
            Yamashita, R., Yamamoto, J., Sekine, M., Tsuritani, K., Wakaguri, H.,
            Ishii, S., Sugiyama, T., Saito, K., Isono, Y., Irie, R., Kushida, N.,
            Yoneyama, T., Otsuka, R., Kanda, K., Yokoi, T., Kondo, H., Wagatsuma, M.,
            Murakawa, K., Ishida, S., Ishibashi, T., Takahashi-Fujii, A.,
            Tanase, T., Nagai, K., Kikuchi, H., Nakai, K., Isogai, T. and Sugano, S.
            Diversification of Transcriptional Modulation: Large-scale
  TITLE
            Identification and Characterization of Putative Alternative
            Promoters of Human Genes
            Genome Res. 16 (1), 55-65 (2006)
  JOURNAL
            16344560
   PUBMED
COMMENT
            Contact: Takao Isogai
            FLJ Project (HRI Team)
            Helix Research Institute
            2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
            Tel: 81-438-52-3975
            Fax: 81-438-52-3986
            Email: flj-cdna@nifty.com
            NEDO human cDNA project (New Energy and Industrial Technology
            Developmental Organization, Japan); cDNA library construction:
            Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
            Research Association for Biotechnology (RAB) and Biotechnology
            Center, National Institute of Technology and Evaluation; 3'-end one
            pass sequencing: RAB.
FEATURES
                     Location/Qualifiers
     source
                     1. .362
                     /organism="Homo sapiens"
                     /mol type="mRNA"
                     /db_xref="taxon:9606"
                     /clone="FCBBF3015237"
                     /tissue_type="brain"
                     /dev stage="fetal"
                     /clone lib="FCBBF3"
                     /note="Vector: pME18SFL3"
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                          16.1%; Score 96.4; DB 9; Length 362;
  Best Local Similarity
                          99.0%;
                                  Pred. No. 1.3e-06;
                                 0: Mismatches
            97; Conservative
                                                   1; Indels
                                                                 0; Gaps
                                                                             0;
            1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qу
              Db
          132 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 73
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RESULT 5

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61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qу
             72 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 35
Db
RESULT 6
AA831793
                                    482 bp
                                             mRNA
LOCUS
           AA831793
                                                     linear
                                                              EST 05-MAR-1998
DEFINITION
           oa59d10.sl NCI CGAP GCB1 Homo sapiens cDNA clone IMAGE:1309267 3'
           similar to SW:SP97 HUMAN Q12959 PRESYNAPTIC PROTEIN SAP97 ;, mRNA
           sequence.
ACCESSION
           AA831793
VERSION
           AA831793.1 GI:2904892
KEYWORDS
           EST.
           Homo sapiens (human)
SOURCE
 ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
              (bases 1 to 482)
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
 TITLE
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
 JOURNAL
           Unpublished (1997)
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
           Ph.D., Gerald Marti, M.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
           Bonaldo, Ph.D.
            cDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           www-bio.llnl.gov/bbrp/image/image.html
           Insert Length: 921
                               Std Error: 0.00
           Seq primer: -40ml3 fwd. ET from Amersham
           High quality sequence stop: 467.
FEATURES
                    Location/Qualifiers
    source
                    1. .482
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                    /mol type="mRNA"
                    /db_xref="taxon:9606"
                    /clone="IMAGE:1309267"
                    /tissue_type="germinal center B cell"
                    /lab host="DH10B"
                    /clone_lib="NCI_CGAP_GCB1"
                    /note="Vector: pT7T3D-PacI; Site_1: Not I; Site_2: Eco RI;
                    1st strand cDNA was prepared from human tonsillar cells
                    enriched for germinal center B cells by flow sorting
                    (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr.
                    David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA
                    synthesis was primed with a Not I - oligo(dT) primer
                    ]. Double-stranded cDNA was ligated to Eco RI adaptors
```

(Pharmacia), digested with Not I and cloned into the Not I

and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization, and was constructed by Bento Soares and M. Fatima Bonaldo."

ORIGIN

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Query Match
                         16.1%;
                                Score 96.4; DB 1; Length 482;
 Best Local Similarity
                         99.0%; Pred. No. 1.3e-06;
 Matches
           97; Conservative
                               0; Mismatches
                                                1;
                                                    Indels
                                                              0;
                                                                  Gaps
                                                                          0;
           1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qy
             21 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 80
Db
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qу
             81 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 118
Db
RESULT 7
BU741256
LOCUS
           BU741256
                                   531 bp
                                             mRNA
                                                    linear
                                                             EST 10-OCT-2002
DEFINITION UI-E-EJ0-air-d-04-0-UI.sl UI-E-EJ0 Homo sapiens cDNA clone
           UI-E-EJ0-air-d-04-0-UI 3', mRNA sequence.
           BU741256
ACCESSION
VERSION
           BU741256.1 GI:23684413
KEYWORDS
           EST.
SOURCE
           Homo sapiens (human)
           Homo sapiens
 ORGANISM
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
              (bases 1 to 531)
REFERENCE
           1
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
 AUTHORS
  TITLE
           Normalization and subtraction: two approaches to facilitate gene
           discovery
  JOURNAL
           Genome Res. 6 (9), 791-806 (1996)
   PUBMED
           8889548
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           Tissue Procurement: Dr. Gregg Hageman
            cDNA Library preparation: Dr. M. Bento Soares, University of Iowa
            cDNA Library Arrayed by: Dr. M. Bento Soares, University of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, University of Iowa
            Clone Distribution: Researchers may obtain clones from Research
           Genetics (www.resgen.com).
           Seq primer: M13 FORWARD
           POLYA=Yes.
FEATURES
                    Location/Qualifiers
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                    /organism="Homo sapiens"
                    /mol type="mRNA"
                    /db xref="taxon:9606"
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optic nerve, retina, Retina Foveal and Macular, RPE and
Choroid"
/dev stage="fetal and adult"
/lab host="DH10B (Life Technologies) (T1 phage resistant)"
/clone lib="UI-E-EJ0"
/note="Organ: eye; Vector: pT7T3-Pac (Pharmacia) with a
modified polylinker; Site_1: EcoR I; Site 2: Not I;
UI-E-EJ0 is a subtracted cDNA library constructed
according to Bonaldo, Lennon and Soares, Genome Research,
6:791-806, 1996. First strand cDNA synthesis was primed
with an oligo-dT primer containing a Not I site. Double
stranded cDNA was ligated to an EcoR I adaptor, digested
with Not I, and cloned directionally into pT7T3-Pac
vector. The oligonucleotide used to prime the synthesis of
first-strand cDNA contains a library tag sequence that is
located between the Not I site and the (dT)18 tail. The
sequence tags for this library are: fetal eyes,
AGAATCAAGA; lens, CGATTAGCGA; eye anterior segment,
AATGCCGCAT; optic nerve, CCATTAAGTG; retina, CCGCG; Retina
Foveal and Macular, GTCC; RPE and Choroid, ACCTA. This
library was created for the program, Gene Discovery in the
Visual System, supported by National Eye Institute (NEI).
TAG TISSUE=human lens
TAG LIB=UI-E-EJ0
TAG SEQ=CGATTAGCGA"
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ORIGIN

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Query Match
                    16.1%;
                          Score 96.4; DB 3; Length 531;
 Best Local Similarity
                    99.0%; Pred. No. 1.3e-06;
 Matches
         97; Conservative
                          0; Mismatches
                                        1;
                                           Indels
                                                             0;
         1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qу
           Db
        30 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 89
        61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qу
           Db
        90 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 127
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RESULT 8
BM727000/c
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LOCUS BM727000 mRNA 532 bp linear EST 01-MAR-2002 DEFINITION

UI-E-EJ0-air-d-04-0-UI.rl UI-E-EJ0 Homo sapiens cDNA clone

UI-E-EJ0-air-d-04-0-UI 5', mRNA sequence.

ACCESSION BM727000

VERSION BM727000.1 GI:19048333

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;

Hominidae: Homo.

(bases 1 to 532) REFERENCE 1

AUTHORS Bonaldo, M.F., Lennon, G. and Soares, M.B.

```
JOURNAL
           Genome Res. 6 (9), 791-806 (1996)
  PUBMED
           8889548
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           Tissue Procurement: Dr. Gregg Hageman
            cDNA Library preparation: Dr. M. Bento Soares, Univeristy of Iowa
            cDNA Library Arrayed by: Dr. M. Bento Soares, Univeristy of Iowa
            DNA Sequencing by: Dr. M. Bento Soares, Univeristy of Iowa
            Clone Distribution: Researchers may obtain clones from Research
           Genetics (www.resgen.com).
           Seq primer: M13 Reverse.
FEATURES
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                    /organism="Homo sapiens"
                    /mol type="mRNA"
                    /db_xref="taxon:9606"
                    /clone="UI-E-EJ0-air-d-04-0-UI"
                    /tissue type="fetal eyes, lens, eye anterior segment,
                    optic nerve, retina, Retina Foveal and Macular, RPE and
                    Choroid"
                    /dev_stage="fetal and adult"
                    /lab host="DH10B (Life Technologies) (T1 phage resistant)"
                    /clone lib="UI-E-EJ0"
                    /note="Organ: eye; Vector: pT7T3-Pac (Pharmacia) with a
                    modified polylinker; Site_1: EcoR I; Site_2: Not I;
                    UI-E-EJO is a subtracted cDNA library constructed
                    according to Bonaldo, Lennon and Soares, Genome Research,
                    6:791-806, 1996. First strand cDNA synthesis was primed
                    with an oligo-dT primer containing a Not I site. Double
                    stranded cDNA was ligated to an EcoR I adaptor, digested
                    with Not I, and cloned directionally into pT7T3-Pac
                    vector. The oligonucleotide used to prime the synthesis of
                    first-strand cDNA contains a library tag sequence that is
                    located between the Not I site and the (dT)18 tail. The
                    sequence tags for this library are: fetal eyes,
                    AGAATCAAGA; lens, CGATTAGCGA; eye anterior segment,
                    AATGCCGCAT; optic nerve, CCATTAAGTG; retina, CCGCG; Retina
                    Foveal and Macular, GTCC; RPE and Choroid, ACCTA. This
                     library was created for the program, Gene Discovery in the
                    Visual System, supported by National Eye Institute (NEI)."
ORIGIN
                                 Score 96.4; DB 3;
                                                    Length 532;
  Query Match
                         16.1%;
  Best Local Similarity
                         99.0%; Pred. No. 1.3e-06;
           97: Conservative
                                0;
                                   Mismatches
  Matches
                                                  1; Indels
                                                                0;
                                                                    Gaps
                                                                            0;
            1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qy
              493 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 434
Db
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Normalization and subtraction: two approaches to facilitate gene

TITLE

discovery

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61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qу
              433 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 396
Db
RESULT 9
DA315109/c
LOCUS
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                                     578 bp
                                               mRNA
                                                       linear
                                                                EST 30-OCT-2005
DEFINITION
           DA315109 BRHIP3 Homo sapiens cDNA clone BRHIP3005378 5', mRNA
            sequence.
ACCESSION
            DA315109
VERSION
            DA315109.1 GI:78306086
KEYWORDS
            EST.
SOURCE
            Homo sapiens (human)
  ORGANISM
            Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE
               (bases 1 to 578)
            1
  AUTHORS
            Kimura, K., Wakamatsu, A., Suzuki, Y., Ota, T., Nishikawa, T.,
            Yamashita, R., Yamamoto, J., Sekine, M., Tsuritani, K., Wakaquri, H.,
            Ishii, S., Sugiyama, T., Saito, K., Isono, Y., Irie, R., Kushida, N.,
            Yoneyama, T., Otsuka, R., Kanda, K., Yokoi, T., Kondo, H., Wagatsuma, M.,
            Murakawa, K., Ishida, S., Ishibashi, T., Takahashi-Fujii, A.,
            Tanase, T., Nagai, K., Kikuchi, H., Nakai, K., Isogai, T. and Sugano, S.
  TITLE
            Diversification of Transcriptional Modulation: Large-scale
            Identification and Characterization of Putative Alternative
            Promoters of Human Genes
  JOURNAL
            Genome Res. 16 (1), 55-65 (2006)
   PUBMED
            16344560
COMMENT
            Contact: Takao Isogai
            FLJ Project (HRI Team)
            Helix Research Institute
            2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
            Tel: 81-438-52-3975
            Fax: 81-438-52-3986
            Email: flj-cdna@nifty.com
            NEDO human cDNA project (New Energy and Industrial Technology
            Developmental Organization, Japan); cDNA library construction:
            Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
            Research Association for Biotechnology (RAB) and Biotechnology
            Center, National Institute of Technology and Evaluation; 3'-end one
            pass sequencing: RAB.
FEATURES
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                          99.0%; Pred. No. 1.3e-06;
            97; Conservative
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                                                                     Gaps
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1 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
Qу
              264 TACAATAGACCCTGCTTCTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 205
Db
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QУ
              Db
         204 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 167
RESULT 10
DA262508/c
LOCUS
           DA262508
                                    580 bp
                                              mRNA
                                                      linear
                                                              EST 03-NOV-2005
DEFINITION DA262508 BRCAN2 Homo sapiens cDNA clone BRCAN2015005 5', mRNA
           sequence.
ACCESSION
           DA262508
           DA262508.1 GI:79168489
VERSION
KEYWORDS
           EST.
SOURCE
           Homo sapiens (human)
 ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
               (bases 1 to 580)
 AUTHORS
           Kimura, K., Wakamatsu, A., Suzuki, Y., Ota, T., Nishikawa, T.,
           Yamashita, R., Yamamoto, J., Sekine, M., Tsuritani, K., Wakaguri, H.,
           Ishii, S., Sugiyama, T., Saito, K., Isono, Y., Irie, R., Kushida, N.,
           Yoneyama, T., Otsuka, R., Kanda, K., Yokoi, T., Kondo, H., Wagatsuma, M.,
           Murakawa, K., Ishida, S., Ishibashi, T., Takahashi-Fujii, A.,
           Tanase, T., Nagai, K., Kikuchi, H., Nakai, K., Isogai, T. and Sugano, S.
 TITLE
           Diversification of Transcriptional Modulation: Large-scale
           Identification and Characterization of Putative Alternative
           Promoters of Human Genes
  JOURNAL
           Genome Res. 16 (1), 55-65 (2006)
  PUBMED
           16344560
COMMENT
           Contact: Takao Isogai
           FLJ Project (HRI Team)
           Helix Research Institute
           2-6-7 Kazusa-Kamatari, Kisarazu, Chiba, 292-0818, Japan
           Tel: 81-438-52-3975
           Fax: 81-438-52-3986
           Email: flj-cdna@nifty.com
           NEDO human cDNA project (New Energy and Industrial Technology
           Developmental Organization, Japan); cDNA library construction:
           Helix Research Institute (HRI); 5'-end one pass sequencing: HRI,
           Research Association for Biotechnology (RAB) and Biotechnology
           Center, National Institute of Technology and Evaluation; 3'-end one
           pass sequencing: RAB.
FEATURES
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Qу
             Db
         218 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 159
          61 ATCTACTTCATTTACTTGTAATATACAGTCATTGACCC 98
Qy
             158 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 121
Db
RESULT 11
BF677732/c
LOCUS
           BF677732
                                   842 bp
                                             mRNA
                                                     linear
                                                             EST 21-DEC-2000
DEFINITION
           602085430F1 NIH MGC 83 Homo sapiens cDNA clone IMAGE:4249854 5',
           mRNA sequence.
           BF677732
ACCESSION
           BF677732.1 GI:11951627
VERSION
KEYWORDS
SOURCE
           Homo sapiens (human)
 ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
              (bases 1 to 842)
 AUTHORS
           NIH-MGC http://mgc.nci.nih.gov/.
           National Institutes of Health, Mammalian Gene Collection (MGC)
 TITLE
  JOURNAL
           Unpublished (1999)
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: CLONETECH Laboratories, Inc.
            cDNA Library Preparation: CLONETECH Laboratories, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           http://image.llnl.gov
           Plate: LLCM1070 row: k column: 07
           High quality sequence stop: 614.
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                    /lab_host="DH10B (T1 phage-resistant)"
                    /clone lib="NIH MGC 83"
                    /note="Organ: prostate; Vector: pDNR-LIB (Clontech);
                    Site_1: SfiI (ggccgcctcggcc); Site_2: SfiI
                    (ggccattatggcc); 5' and 3' adaptors were used in cloning
                    as follows: 5' adaptor sequence: 5'-CACGGCCATTATGGCC-3'
                    and 3' adaptor sequence:
                    5'-ATTCTAGAGGCCGAGGCGGCCGACATG-dT(30)BN-3' (where B = A,
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C, or G and N = A, C, G, or T). Average insert size 1.4 kb (range 0.5-4.0 kb). 14/15 colonies contained inserts by PCR. This library was enriched for full-length clones and was constructed by Clontech Laboratories (Palo Alto, CA)."
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ORIGIN

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Query Match
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 Best Local Similarity
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 Matches
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                                                    Indels
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                                                                 Gaps
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Qy
           1 TACAATAGACCCTGCTTCTTTCAACGCTTCAACTGCTTTGCTATGTGTTACATCACGAAC 60
             Db
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Qу
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             Db
         243 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 206
RESULT 12
BG164149/c
LOCUS
           BG164149
                                   860 bp
                                            mRNA
                                                    linear EST 06-FEB-2001
DEFINITION
           602343020F1 NIH_MGC_89 Homo sapiens cDNA clone IMAGE:4453276 5',
           mRNA sequence.
ACCESSION
           BG164149
VERSION
           BG164149.1 GI:12670852
KEYWORDS
           EST.
SOURCE
           Homo sapiens (human)
 ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
              (bases 1 to 860)
 AUTHORS
           NIH-MGC http://mgc.nci.nih.gov/.
 TITLE
           National Institutes of Health, Mammalian Gene Collection (MGC)
 JOURNAL
           Unpublished (1999)
           Contact: Robert Strausberg, Ph.D.
COMMENT
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: ATCC
            cDNA Library Preparation: Life Technologies, Inc.
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)
            DNA Sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           http://image.llnl.gov
           Plate: LLAM10243 row: g column: 05
           High quality sequence stop: 687.
FEATURES
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Site 2: SalI; Cloned unidirectionally; oligo-dT primed.
                    Average insert size 1.3 kb. Library enriched for
                    full-length clones and constructed by Life Technologies.
                    Note: this is a NIH MGC Library."
                                 Score 96.4; DB 2; Length 860;
 Query Match
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                                 Pred. No. 1.2e-06;
 Best Local Similarity
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           97; Conservative
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                                                     Indels
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                                                                  Gaps
                                                                           0;
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           AY420526
                                   2675 bp
                                             DNA
                                                     linear
                                                              GSS 17-DEC-2003
DEFINITION Pan troglodytes DLG1 gene, VIRTUAL TRANSCRIPT, partial sequence,
           genomic survey sequence.
           AY420526
           AY420526.1 GI:39776483
           GSS.
           Pan troglodytes (chimpanzee)
           Pan troglodytes
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Pan.
              (bases 1 to 2675)
           Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
           Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
           Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
           Adams, M.D. and Cargill, M.
           Inferring nonneutral evolution from human-chimp-mouse orthologous
           gene trios
           Science 302 (5652), 1960-1963 (2003)
           14671302
               (bases 1 to 2675)
           2
           Clark, A.G., Glanowski, S., Nielson, R., Thomas, P., Kejariwal, A.,
           Todd, M.A., Tanenbaum, D.M., Civello, D.R., Lu, F., Murphy, B.,
           Ferriera, S., Wang, G., Zheng, X.H., White, T.J., Sninsky, J.J.,
           Adams, M.D. and Cargill, M.
           Direct Submission
           Submitted (16-NOV-2003) Celera Genomics, 45 West Gude Drive,
           Rockville, MD 20850, USA
           This sequence as made by sequencing genomic exons and ordering them
           based on alignment.
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                    1. .2675
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                    /mol type="genomic DNA"
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/note="Organ: kidney; Vector: pCMV-SPORT6; Site_1: NotI;

ORIGIN

Qу

Db

Qу

Db

RESULT 13 AY420526/c LOCUS

ACCESSION

REFERENCE

TITLE

JOURNAL

AUTHORS

REFERENCE

TITLE JOURNAL

COMMENT

FEATURES

source

/db_xref="taxon:9598"

PUBMED

AUTHORS

ORGANISM

VERSION **KEYWORDS**

SOURCE

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                                                                  Gaps
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Db
RESULT 14
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VERSION
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SOURCE
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           Hominidae; Homo.
REFERENCE
              (bases 1 to 447)
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 AUTHORS
  TITLE
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
 JOURNAL
           Unpublished (1997)
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
           Ph.D., Gerald Marti, M.D.
            cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
           Bonaldo, Ph.D.
            cDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           www-bio.llnl.gov/bbrp/image/image.html
           Possible reversed clone: similarity on wrong strand
           Seq primer: -28m13 rev1 ET from Amersham
           High quality sequence stop: 385.
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<1. .>2675

gene

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                 /clone="IMAGE:814570"
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                 /lab host="DH10B"
                  /clone lib="NCI CGAP GCB1"
                 /note="Vector: pT7T3D-PacI; Site 1: Not I; Site 2: Eco RI;
                 1st strand cDNA was prepared from human tonsillar cells
                 enriched for germinal center B cells by flow sorting
                 (CD20+, IgD-), provided by Dr. Louis M. Staudt (NCI), Dr.
                 David Allman (NCI) and Dr. Gerald Marti (CBER). cDNA
                 synthesis was primed with a Not I - oligo(dT) primer
                  ]. Double-stranded cDNA was ligated to Eco RI adaptors
                  (Pharmacia), digested with Not I and cloned into the Not I
                 and Eco RI sites of the modified pT7T3 vector. Library
                 went through one round of normalization, and was
                 constructed by Bento Soares and M. Fatima Bonaldo."
                      15.8%; Score 94.8; DB 1; Length 447;
Best Local Similarity
                      98.0%; Pred. No. 2.5e-06;
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                                                 Indels
                                                           0; Gaps
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           101 ATCTACTTCATTTACTCGTAATATACAGTCATTGACCC 138
         BP229026
                                582 bp
                                          mRNA
                                                 linear
                                                          EST 15-SEP-2004
         BP229026 Sugano cDNA library, fetal brain Homo sapiens cDNA clone
         FBR01745, mRNA sequence.
         BP229026
         BP229026.1 GI:52101936
         Homo sapiens (human)
         Homo sapiens
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         Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
         Hominidae; Homo.
            (bases 1 to 582)
         Suzuki, Y., Yamashita, R., Shirota, M., Sakakibara, Y., Chiba, J.,
         Mizushima-Sugano, J., Nakai, K. and Sugano, S.
         Sequence comparison of human and mouse genes reveals a homologous
         block structure in the promoter regions
         Genome Res. 14 (9), 1711-1718 (2004)
         15342556
         Contact: Yutaka Suzuki
         Department of Virology
         Institute of Medical Science, University of Tokyo
         4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
```

ORIGIN

Qу

Db

Qy

Db

RESULT 15 BP229026/c LOCUS

DEFINITION

ACCESSION

REFERENCE

TITLE

COMMENT

AUTHORS

JOURNAL

PUBMED

ORGANISM

EST.

Tel: 81-3-5449-5343

VERSION KEYWORDS

SOURCE

Query Match

Matches